

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:sssptaul532cxa

PASSWORD:

LOGINID/PASSWORD REJECTED

The loginid and/or password sent to STN were invalid.  
You either typed them incorrectly, or line noise may  
have corrupted them.

Do you wish to retry the logon?

Enter choice (y/N):

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LOGINID:sssptaul53cxa

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

\* \* \* \* \* Welcome to STN International \* \* \* \* \*

NEWS	1	Web Page URLs for STN Seminar Schedule - N. America
NEWS	2	"Ask CAS" for self-help around the clock
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NEWS	5	NOV 30 PHAR reloaded with additional data
NEWS	6	DEC 01 LISA now available on STN
NEWS	7	DEC 09 12 databases to be removed from STN on December 31, 2004
NEWS	8	DEC 15 MEDLINE update schedule for December 2004
NEWS	9	DEC 17 ELCOM reloaded; updating to resume; current-awareness alerts (SDIs) affected

NEWS 10 DEC 17 COMPUAB reloaded; updating to resume; current-awareness alerts (SDIs) affected

NEWS 11 DEC 17 SOLIDSTATE reloaded; updating to resume; current-awareness alerts (SDIs) affected

NEWS 12 DEC 17 CERAB reloaded; updating to resume; current-awareness alerts (SDIs) affected

NEWS 13 DEC 17 THREE NEW FIELDS ADDED TO IFIPAT/IFIUDB/IFICDB

NEWS 14 DEC 30 EPFULL: New patent full text database to be available on STN

NEWS 15 DEC 30 CAPLUS - PATENT COVERAGE EXPANDED

NEWS 16 JAN 03 No connect-hour charges in EPFULL during January and February 2005

NEWS 17 JAN 26 CA/CAPLUS - Expanded patent coverage to include the Russian Agency for Patents and Trademarks (ROSPATENT)

NEWS 18 FEB 10 STN Patent Forums to be held in March 2005

NEWS EXPRESS JANUARY 10 CURRENT WINDOWS VERSION IS V7.01a, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 10 JANUARY 2005

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\* \* \* \* \* STN Columbus \* \* \* \* \*

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=> file caplus uspatful japio medline biosis embase

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=> s (civamide or (vanillyl(w)6(w)nonenamide))  
 L1 352 (CIVAMIDE OR (VANILLYL(W) 6(W) NONENAMIDE))

=> s l1 an (headache or neuralgia or neuropathy)  
MISSING OPERATOR L1 AN  
The search profile that was entered contains terms or  
nested terms that are not separated by a logical operator.

=> s l1 and (headache or neuralgia or neuropathy)  
L2 59 L1 AND (HEADACHE OR NEURALGIA OR NEUROPATHY)

=> s l2 and (topical? or intranasal? or nasal?)  
L3 49 L2 AND (TOPICAL? OR INTRANASAL? OR NASAL?)

=> s l3 and (drug delivery)  
1 FILES SEARCHED...  
L4 6 L3 AND (DRUG DELIVERY)

=> d l4 1-6 ibib abs

L4 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN  
ACCESSION NUMBER: 2004:805904 CAPLUS  
DOCUMENT NUMBER: 142:85673  
TITLE: **Intranasal** medications for the treatment of  
migraine and cluster **headache**  
AUTHOR(S): Rapoport, Alan M.; Bigal, Marcelo E.; Tepper, Stewart  
J.; Sheftell, Fred D.  
CORPORATE SOURCE: Columbia University College of Physicians & Surgeons,  
New York, NY, USA  
SOURCE: CNS Drugs (2004), 18(10), 671-685  
CODEN: CNDREF; ISSN: 1172-7047  
PUBLISHER: Adis International Ltd.  
DOCUMENT TYPE: Journal; General Review  
LANGUAGE: English

AB A review. **Intranasal** medications for the treatment of  
**headache** have recently received increased attention. This paper  
reviews **intranasal** formulations of a variety of available  
medications (dihydroergotamine mesylate, sumatriptan, zolmitriptan,  
butorphanol, capsaicin and lidocaine) and one exptl. medication (  
**civamide**, a cis-isomer of capsaicin) for the treatment of migraine  
and cluster **headache**. Although the efficacy of  
**intranasal** agents varies with the product used, **intranasal**  
delivery may be both convenient and more effective than other modes of  
**drug delivery** for a variety of reasons: (i)  
**intranasal** administration bypasses small bowel gastrointestinal  
tract absorption, which is often significantly delayed during the acute  
phase of a migraine attack; (ii) nauseated patients may prefer non-oral  
formulations as they decrease the chance of vomiting and are more rapidly  
effective; (iii) **intranasal** administration causes no pain or  
injection site reaction and is easier and more convenient to administer  
than injection or suppository and so may be used earlier in a migraine  
attack, resulting in better efficacy; (iv) **intranasal** medication  
produces the same number or fewer adverse events than injections; and (v)  
**intranasal** formulations offer a more rapid onset of action than  
oral medications, for some of the above reasons and, as such, may be more  
useful in patients with cluster **headache**, although this needs to  
be verified. However, it is important to emphasize that a preference  
study showed that most patients prefer oral tablets to an  
**intranasal** formulation. Also, some **nasal** preps. have  
significant adverse effects or are not well absorbed and therefore do not  
work consistently; others are more challenging to administer as a result  
of their delivery apparatus. Nevertheless, it is our opinion that **nasal**  
preps. increase therapeutic options and may result in faster response  
times and better efficacy than oral formulations and better patient  
satisfaction than injectable preps.

REFERENCE COUNT: 80 THERE ARE 80 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 6 USPATFULL on STN

ACCESSION NUMBER: 2000:61198 USPATFULL  
TITLE: Therapeutic uses of pungent botanicals and their  
related compounds  
INVENTOR(S): Staggs, Jeff J., 7474 E. Arkansas Ave. #8-10, Denver,  
CO, United States 80231

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6063381		20000516
	WO 9323061		19931125
APPLICATION INFO.:	US 1997-338489		19970318 (8)
	WO 1993-US4763		19930519
			19970318 PCT 371 date
			19970318 PCT 102(e) date
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Weddington, Kevin E.		
NUMBER OF CLAIMS:	19		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	13 Drawing Figure(s); 7 Drawing Page(s)		
LINE COUNT:	2066		

AB A new class of general antiinfective agents extracted from pepper, ginger, and other plant species containing vanillyl and piperidine ring structures typical of the pungent principal found in pepper and ginger. The role of these structures, their attached hydrocarbons groups, and other agents found with the plant extract is demonstrated in the topical treatment of dermatophyte infections, tissue injuries, and abnormal proliferations of keratin.

L4 ANSWER 3 OF 6 USPATFULL on STN

ACCESSION NUMBER: 1998:64759 USPATFULL  
TITLE: Method and compositions for controlling oral and  
pharyngeal pain using capsaicinoids  
INVENTOR(S): Byas-Smith, Michael G., Decatur, GA, United States  
PATENT ASSIGNEE(S): Emory University, Atlanta, GA, United States (U.S.  
corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5762963		19980609
APPLICATION INFO.:	US 1995-478554		19950607 (8)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Sayala, Chhaya D.		
LEGAL REPRESENTATIVE:	Knowles, Sherry M. King & Spalding		
NUMBER OF CLAIMS:	45		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 1 Drawing Page(s)		
LINE COUNT:	1234		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Methods and compositions are provided for the oral delivery of temporally increasing concentrations of capsaicin, its derivatives, and analogs (collectively, "capsaicinoids"), to provide oral or pharyngeal analgesia while minimizing sensations of nausea and burning associated with the oral administration of capsaicinoids. The methods and compositions described herein soothe and relieve oral or pharynx pain. In one embodiment, one or more capsaicinoids are dispersed within a lollipop, with successively decreasing concentrations of capsaicin from the center out to the periphery, and administered to a patient in need of amelioration of oral pain. Alternatively, the capsaicinoid can be dispersed, with decreasing concentrations from the center to the periphery, in a tablet, caplet, lozenge, troche, pill, candy, or similar formulation. Capsaicinoids include dihydrocapsaicin, norhydrocapsaicin,

homocapsaicin, homodihydrocapsaicin I, norhydrocapsaicin, homodihydrocapsaicin, nordihydrocapsaicin, **civamide**, nonivamide, NE-19550 (also called olvanil), NE-21610, NE-28345 (also called N-oley-l-homovanillamide), their analogs, and derivatives.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L4 ANSWER 4 OF 6 USPATFULL on STN

ACCESSION NUMBER: 97:80931 USPATFULL  
TITLE: Transdermal therapeutic formulation  
INVENTOR(S): Davis, Roosevelt, 27 Lullwater Estate Rd., Atlanta, GA, United States 30307  
Primo-Davis, Susan A., 27 Lullwater Estate Rd., Atlanta, GA, United States 30307

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5665378		19970909
APPLICATION INFO.:	US 1995-560806		19951121 (8)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1994-315343, filed on 30 Sep 1994, now abandoned		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Phelan, D. Gabrielle		
LEGAL REPRESENTATIVE:	Connolly & Hutz		
NUMBER OF CLAIMS:	19		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	1 Drawing Figure(s); 1 Drawing Page(s)		
LINE COUNT:	473		

AB The present invention relates to a transdermal therapeutic formulation comprising capsaicin, a nonsteroidal anti-inflammatant and pamabrom. The formulation is used to alleviate pain or discomfort in a mammal by being applied to the skin of the mammal thereby causing the active ingredients in the formulation to pass into and/or through the skin of the mammal. In a preferred embodiment of the present invention, the formulation is used in patch form for the treatment of the pain and discomfort associated with menstrual cramps, water retention (e.g., "bloating") and/or muscular pain (e.g., muscular back pain).

L4 ANSWER 5 OF 6 MEDLINE on STN

ACCESSION NUMBER: 2004368141 MEDLINE  
DOCUMENT NUMBER: PubMed ID: 15270595  
TITLE: Intranasal medications for the treatment of migraine and cluster headache.  
AUTHOR: Rapoport Alan M; Bigal Marcelo E; Tepper Stewart J; Sheftell Fred D  
CORPORATE SOURCE: Columbia University College of Physicians & Surgeons, New York, NY, USA.. alanrapoport@nech.net  
SOURCE: CNS drugs, (2004) 18 (10) 671-85. Ref: 80  
Journal code: 9431220. ISSN: 1172-7047.  
PUB. COUNTRY: New Zealand  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, TUTORIAL)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 200410  
ENTRY DATE: Entered STN: 20040725  
Last Updated on STN: 20041026  
Entered Medline: 20041025

AB Intranasal medications for the treatment of headache have recently received increased attention. This paper reviews intranasal formulations of a variety of available medications (dihydroergotamine mesylate [dihydroergotamine mesilate], sumatriptan,

zolmitriptan, butorphanol, capsaicin and lidocaine [lignocaine]) and one experimental medication (**civamide**, a cis-isomer of capsaicin) for the treatment of migraine and cluster **headache**. Although the efficacy of **intranasal** agents varies with the product used, **intranasal** delivery may be both convenient and more effective than other modes of **drug delivery** for a variety of reasons:

(i) **intranasal** administration bypasses small bowel gastrointestinal tract absorption, which is often significantly delayed during the acute phase of a migraine attack; (ii) nauseated patients may prefer non-oral formulations as they decrease the chance of vomiting and are more rapidly effective; (iii) **intranasal** administration causes no pain or injection site reaction and is easier and more convenient to administer than injection or suppository and so may be used earlier in a migraine attack, resulting in better efficacy; (iv) **intranasal** medication produces the same number or fewer adverse events than injections; and (v) **intranasal** formulations offer a more rapid onset of action than oral medications, for some of the above reasons and, as such, may be more useful in patients with cluster **headache**, although this needs to be verified. However, it is important to emphasise that a preference study showed that most patients prefer oral tablets to an **intranasal** formulation. Also, some **nasal** preparations have significant adverse effects or are not well absorbed and therefore do not work consistently; others are more challenging to administer as a result of their delivery apparatus. Nevertheless, it is our opinion that **nasal** preparations increase therapeutic options and may result in faster response times and better efficacy than oral formulations and better patient satisfaction than injectable preparations.

L4 ANSWER 6 OF 6 EMBASE COPYRIGHT 2005 ELSEVIER INC. ALL RIGHTS RESERVED.  
on STN

ACCESSION NUMBER: 2004369350 EMBASE

TITLE: **Intranasal** medications for the treatment of migraine and cluster **headache**.

AUTHOR: Rapoport A.M.; Bigal M.E.; Tepper S.J.; Sheftell F.D.

CORPORATE SOURCE: Dr. A.M. Rapoport, New England Center for Headache, P.C., 778 Long Ridge Road, Stamford, CT 06902-1251, United States. alanrapoport@nech.net

SOURCE: CNS Drugs, (2004) 18/10 (671-685).

Refs: 80

ISSN: 1172-7047 CODEN: CNDREF

COUNTRY: New Zealand

DOCUMENT TYPE: Journal; General Review

FILE SEGMENT: 008 Neurology and Neurosurgery

030 Pharmacology

037 Drug Literature Index

038 Adverse Reactions Titles

039 Pharmacy

LANGUAGE: English

SUMMARY LANGUAGE: English

AB **Intranasal** medications for the treatment of **headache**

have recently received increased attention. This paper reviews **intranasal** formulations of a variety of available medications (dihydroergotamine mesylate [dihydroergotamine mesilate], sumatriptan, zolmitriptan, butorphanol, capsaicin and lidocaine [lignocaine]) and one experimental medication (**civamide**, a cis-isomer of capsaicin) for the treatment of migraine and cluster **headache**. Although the efficacy of **intranasal** agents varies with the product used, **intranasal** delivery may be both convenient and more effective than other modes of **drug delivery** for a variety of reasons:

(i) **intranasal** administration bypasses small bowel gastrointestinal tract absorption, which is often significantly delayed during the acute phase of a migraine attack; (ii) nauseated patients may prefer non-oral formulations as they decrease the chance of vomiting and are more rapidly effective; (iii) **intranasal** administration

causes no pain or injection site reaction and is easier and more convenient to administer than injection or suppository and so may be used earlier in a migraine attack, resulting in better efficacy; (iv) **intranasal** medication produces the same number or fewer adverse events than injections; and (v) **intranasal** formulations offer a more rapid onset of action than oral medications, for some of the above reasons and, as such, may be more useful in patients with cluster headache, although this needs to be verified. However, it is important to emphasise that a preference study showed that most patients prefer oral tablets to an **intranasal** formulation. Also, some **nasal** preparations have significant adverse effects or are not well absorbed and therefore do not work consistently; others are more challenging to administer as a result of their delivery apparatus. Nevertheless, it is our opinion that **nasal** preparations increase therapeutic options and may result in faster response times and better efficacy than oral formulations and better patient satisfaction than injectable preparations.

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